IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of: GARDELLA *et al.*	Confirmation No.: 2201
Appl. No.: 10/549,592 (U.S. Nat'l Phase of PCT/US2003/008261)	Art Unit: 1654
§ 371 Date: January 12, 2007	Examiner: *To Be Assigned*
For: **Conformationally Constrained Parathyroid Hormones With α-Helix Stabilizers**	Atty. Docket: 0609.5150000/TJS/PAC

Information Disclosure Statement under 37 C.F.R. § 1.97(b)

Mail Stop Amendment

Commissioner for Patents
PO Box 1450
Alexandria, VA 22313-1450

Sir:

Listed on accompanying IDS Forms, PTO/SB/08A and PTO/SB/08B are documents that may be considered material to the examination of this application, in compliance with the duty of disclosure requirements of 37 C.F.R. §§ 1.56, 1.97 and 1.98.

Copies of documents **FP1** and **NPL1-NPL36** are submitted. However, in accordance with 37 C.F.R. § 1.98(a)(2), copies of U.S. patents, documents **US1** to **US5,** cited on the attached IDS Form, PTO/SB/08A, are not submitted.

Where the publication date of a listed document does not provide a month of publication, the year of publication of the listed document is sufficiently earlier than the effective U.S. filing date and any foreign priority date so that the month of publication is not in issue. Applicants have listed publication dates on the attached IDS Forms based on information presently available to the undersigned. However, the listed publication dates should not be construed as an admission that the information was actually published on the date indicated.

Applicants reserve the right to establish the patentability of the claimed invention over any of the information provided herewith, and/or to prove that this information may

not be prior art, and/or to prove that this information may not be enabling for the teachings purportedly offered.

This statement should not be construed as a representation that a search has been made, or that information more material to the examination of the present patent application does not exist. The Examiner is specifically requested not to rely solely on the material submitted herewith.

This Information Disclosure Statement is being filed before the mailing date of a first Office Action on the merits. No statement or fee is required.

It is respectfully requested that the Examiner initial and return a copy of the enclosed IDS Forms, and indicate in the official file wrapper of this patent application that the documents have been considered.

The U.S. Patent and Trademark Office is hereby authorized to charge any fee deficiency, or credit any overpayment, to our Deposit Account No. 19-0036.

Respectfully submitted,

STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.

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Sheet 1 of 1

Complete if Known	
Application Number	10/549,592 (U.S. Nat'l Phase of PCT/US2003/008261)
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First Named Inventor	GARDELLA, Thomas J.
Art Unit	1654
Examiner Name	*To Be Assigned*
Attorney Docket Number	0609.5150000/TJS/PAC

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Document Number Number-Kind Code[2] (If Known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
/R.T./	US1	4,675,189	06/23/1987	Kent *et al.*	
	US2	4,698,328	10/06/1987	Neer *et al.*	
	US3	4,761,406	08/02/1988	Flora *et al.*	
	US4	6,030,709	02/29/2000	Adermann *et al.*	
	US5	2002/0110871	08/15/2002	Zahradnik *et al.*	

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Foreign Patent Document Country Code[3] Number[4] Kind Code[5] (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T[6]
/R.T./	FP1	WO 97/02834	01/30/1997	Biomeasure Incorporated		

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Examiner Signature	/Roy Teller/	Date Considered	10/09/2009

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/R.T./	NPL1	Barden, J.A. and Kemp, B.E., "NMR Solution Structure of Human Parathyroid Hormone(1-34)," *Biochemistry 32*:7126-7132, American Chemical Society (1993)	
	NPL2	Behar, V., *et al.*, "Photoaffinity Cross-linking Identifies Differences in the Interactions of an Agonist and an Antagonist with the Parathyroid Hormone/Parathyroid Hormone-related Protein Receptor," *J. Biol. Chem. 275*:9-17, American Society for Biochemistry and Molecular Biology, Inc. (2000)	
	NPL3	Bergwitz, C., *et al.*, "Full Activation of Chimeric Receptors by Hybrids between Parathyroid Hormone and Calcitonin," *J. Biol. Chem. 271*:26469-26472, The American Society for Biochemistry and Molecular Biology, Inc. (1996)	
	NPL4	Berridge, M.J., *et al.*, "Changes in the levels of inositol phosphates after agonist-dependent hydrolysis of membrane phosphoinositides," *Biochem. J. 212*:473-482, The Biochemical Society (1983)	
	NPL5	Bowen, W.P. and Jerman, J.C., "Nonlinear regression using spreadsheets," *Trends Pharmacol Sci 16*:413-417, Elsevier, Science Ltd. (1995)	
	NPL6	Carter, P.H., *et al.*, "Studies of the N-Terminal Region of a Parathyroid Hormone-Related Peptide(1-36) Analog: Receptor Subtype-Selective Agonists, Antagonists, and Photochemical Cross-Linking Agents," *Endocrinol. 140*:4972-4981, The Endocrine Society (1999)	
	NPL7	Chen, Z., *et al.*, "Solution Structure of the Osteogenic 1-31 Fragment of Human Parathyroid Hormone," *Biochemistry 39*:12766-12777, American Chemical Society (2000)	
	NPL8	Chorev, M., *et al.*, "Modifications of Position 12 in Parathyroid Hormone and Parathyroid Hormone Related Protein: Toward the Design of Highly Potent Antagonists," Biochemistry 29:1580-1586, American Chemical Society (1990)	
	NPL9	Dempster, D.W., *et al.*, "Anabolic Actions of Parathyroid Hormone on Bone," *Endocrine Rev. 14*:690-709, The Endocrine Society (1993)	
	NPL10	Dempster, D.W., *et al.*, "Erratum: Anabolic Actions of Parathyroid Hormone on Bone," *Endocrine Rev. 15*:261, The Endocrine Society (1994)	

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/R.T./	NPL11	Fairwell, T., *et al.*, "Total Solid-Phase Synthesis, Purification, and Characterization of Human Parathyroid Hormone-(1-84)," *Biochemistry* 22:2691-2697, American Chemical Society (1983)	
	NPL12	Gronwald, W., *et al.*, "Structure of Recombinant Human Parathyroid Hormone in Solution Using Multidimensional NMR Spectroscopy," *Chem. Hoppe-Seyler 377*:175-186, Walter de Gruyter & Co. (1996)	
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	NPL14	Hoare, S.R.J., *et al.*, "Evaluating the Signal Transduction Mechanism of the Parathyroid Hormone 1 Receptor," *J. Biol. Chem. 276*:7741-7753, American Society for Biochemistry and Molecular Biology, Inc. (2001)	
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	NPL16	Kaul, R and Balram, P., "Stereochemical Control of Peptide Folding," *Bioorg. Med. Chem. 7*:105-117, Elsevier Science Ltd. (1999)	
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	NPL18	Luck, M.D., *et al.*, "The (1-14) Fragment of Parathyroid Hormone (PTH) Activates Intact and Amino-Terminally Truncated PTH-1 Receptors," *Mol. Endocrinol. 13*:670-680, The Endocrine Society (1999)	
	NPL19	Marx, U.C., *et al.*, "Structure of Human Parathyroid Hormone 1-37 in Solution," *J. Biol. Chem. 270*:15194-15202, The American Society for Biochemistry and Molecular Biology, Inc. (1995)	

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/R.T./	NPL20	Marx, U.C., *et al.*, "Structure-Activity Relation of NH_2-terminal Human Parathyroid Hormone Fragments," *J. Biol. Chem 273*:4308-4316, American Society for Biochemistry and Molecular Biology, Inc. (1998)	
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	NPL22	Neer, R.M., *et al.*, "Effect of Parathyroid Hormone (1-34) On Fractures and Bone Mineral Density in Postmenopausal Women with Osteoporosis," *N. Eng. J. Med. 344*:1434-1441, Massachusetts Medical Society (2001)	
	NPL23	Pellegrini, M., *et al.*, "Binding Domain of Human Parathyroid Hormone Receptor: From Conformation to Function," *Biochemistry 37*:12737-12743, American Chemical Society (1998)	
	NPL24	Robinson J.R. ed., "Methods to Achieve Controlled Drug Delivery," in: *Sustained and Controlled Release Drug Delivery Systems*, Marcel Dekker, New York,NY, pp 557-593 (1978)	
	NPL25	Shen, V., *et al.*, "Effects of Combined and Separate Intermittent Administration of Low-Dose Human Parathyroid Hormone Fragment (1-34) and 17β-Estradiol on Bone Histomorphometry in Ovariectomized Rats with Established Osteopenia," *Calcif. Tissue Intl. 50:214-220*, Springer-Verlag Inc. (1992)	
	NPL26	Shimizu, M., *et al.*, "Autoactivation of Type-1 Parathyroid Hormone Receptors Containing a Tethered Ligand," *J. Biol. Chem. 275*:19456-19460, The American Society for Biochemistry and Molecular Biology, Inc. (2000)	
	NPL27	Shimizu, M., *et al.*, "Minimization of Parathyroid Hormone," *J. Biol. Chem. 275*:21836-21843, The American Society for Biochemistry and Molecular Biology, Inc. (2000)	
	NPL28	Shimizu, M., *et al.*, "Enhanced Activity in Parathyroid Hormone-(1-14) and -(1-11): Novel Peptides for Probing Ligand-Receptor Interactions," *Endocrinol. 142*:3068-3074, Endocrine Society (2001)	

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/R.T./	NPL29	Shimizu, N., *et al.*, "Parathyroid Hormone (PTH)-(1-14) and -(1-11) Analogs Conformationally Constrained by α-Aminosobutyric Acid Mediate Full Agonist Responses via the Juxtamembrane Region of the PTH-1 Receptor," *J. Biol. Chem. 276*:49003-49012, The American Society for Biochemistry and Molecular Biology, Inc. (2001)	
	NPL30	Slovik, D.M., *et al.*, "Restoration of Spinal Bone in Osteoporotic Men by Treatment with Human Parathyroid Hormone (1-34) and 1,25-Dihydroxyvitamin D," *J. Bone Min. Res. 1*:377-381, Mary Ann Liebert, Inc. (1986)	
	NPL31	Takasu, H., *et al.*, "Amino Terminal Modifications of Human Parathyroid Hormone (PTH) Selectively Alter Phospholipase C Signaling via the Type 1 PTH Receptor: Implications for Design for Signal-Specific PTH Ligands," *Biochemistry 38*:13453-13460, American Chemical Society (1999)	
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	NPL35	Whitfield, J.F., *et al.*, "Comparison of the Ability of Recombinant Human Parathyroid Hormone, rhPTH-(1-84), and hPTH-(1-31)NH_2 Stimulate Femoral Trabecular Bone Growth in Ovariectomized Rats," *Calcif. Tissue Int. 60*:26-29, Springer-Verlag Inc. (1997)	
	NPL36	Wold, F., "Posttranslational Protein Modifications: Perspectives and Prospects," in *Posttranslational Covalent Modifications of Proteins*, B.C. Johnson, eds., Academic Press, Inc., New York, pp. 1-12 (1983)	

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